



## Donkey anti-Rat IgG (H+L) Cross-Adsorbed Secondary Antibody, DyLight<sup>™</sup> 594

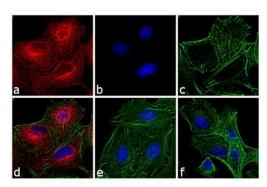
<b>Product Details</b>		
Size	500 μg	
Species Reactivity	Rat	
Host/Isotype	Donkey / IgG	
Class	Polyclonal	
Туре	Secondary Antibody	
Conjugate	DyLight™ 594	
Excitation/Emission Max	587/614 nm	
Immunogen	Rat IgG-heavy and light chain	
Form	Liquid	
Concentration	0.5 mg/mL	
Purification	Antigen affinity chromatography	
Storage buffer	PBS, pH 6.8 to 7.4, with 0.2% BSA	
Contains	0.09% sodium azide	
Storage conditions	4° C	
RRID	AB_2556608	

Applications	Tested Dilution	Publications
Western Blot (WB)	1:5,000-1:20,000	-
Immunohistochemistry (IHC)	1:50-1:2,000	-
Immunocytochemistry (ICC/IF)	1:50-1:500	-
Flow Cytometry (Flow)	1:50-1:200	-
Immunoprecipitation (IP)	Assay-dependent	-

## **Product Specific Information**

This antibody is cross-adsorbed and exhibits minimum reactivity to bovine, chicken, goat, human, mouse, rabbit and sheep.

## Product Images For Donkey anti-Rat IgG (H+L) Cross-Adsorbed Secondary Antibody, DyLight™ 594



Rat IgG (H+L) Cross-Adsorbed Secondary Antibody (SA5-10028) in ICC/IF Immunofluorescence analysis of Donkey anti-Rat IgG (H+L) Cross Adsorbed Secondary Antibody, DyLight 594 conjugate was performed using A549 cells stained with alpha Tubulin (YL1/2) Rat Monoclonal Antibody (Product # MA1-80017). The cells were fixed with 4% paraformaldehyde for 10 minutes, permeabilized with 0.1% Triton™ X-100 for 10 minutes, blocked with 1% BSA for 1 hour and labeled with 2µg/mL Rat primary antibody for 3 hours at room temperature. Donkey anti-Rat IgG (H+L) Cross Adsorbed Secondary Antibody, DyLight 594 conjugate (Product # SA5-10028) was used at a concentration of 1µg /mL in phosphate buffered saline containing 0.2 % BSA for 45 minutes at room temperature, for detection of alpha Tubulin in the cytoplasm (Panel a: red). Nuclei (Panel b: blue) were stained with DAPI in SlowFade® Gold Antifade Mountant (Product # S36938). F-actin was stained with Alexa Fluor® 488 Phalloidin (Product # A12379), 1:300) (Panel c: green). Panel d represents the composite image. No nonspecific staining was observed with the secondary antibody alone (panel f), or with an isotype control (panel e). The images were captured at 60X magnification.

## **□7** References

Three-Dimensional Imaging of Macrophages in Complete Organs. Methods Mol Biol (2023)

Active APPL1 sequestration by Plasmodium favors liver-stage development. Cell Rep (2022)

An In Vivo CRISPR Screen Identifies Stepwise Genetic Dependencies of Metastatic Progression. Cancer Res (2022)

Endothelial and systemic upregulation of miR-34a-5p fine-tunes senescence in progeria. Aging (Albany NY) (2022)

FAM3D is essential for colon homeostasis and host defense against inflammation associated carcinogenesis. Nat Commun (2020)

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